

Claims

1. A method of analysis of amino acids, peptides or proteins, the method comprising:

(1) derivatizing a mixture of amino acids, peptides or proteins, to form at least one amino acid, peptide or protein derivatized to contain a fixed-charge ion, other than at the C-terminal or N-terminal end thereof;

5 (2) introducing the mixture of amino acids, peptides or proteins containing at least one amino acid, peptide or protein derivatized to contain a fixed-charge ion other than at the C-terminal or N-terminal end thereof, to a mass spectrometer;

(3) passing the mixture of amino acids, peptides or proteins containing at least one amino acid, peptide or protein derivatized to contain a fixed-charge ion, other than at the C-terminal or N-terminal end thereof, through a first mass resolving spectrometer to select
10 precursor protein or peptide ions having a first mass-to-charge ratio;

(4) subjecting the precursor ions of the first mass-to-charge ratio to dissociation to form product ions having a second mass-to-charge ratio that is characteristic of a fragmentation occurring at a site adjacent to the fixed charge; and

15 (5) detecting the product ions having the second mass-to-charge ratio.

2. The method of claim 1, wherein the product ions having the second mass-to-charge ratio are product ions formed by neutral loss of the fixed charge from the precursor ions.

20 3. The method of claim 1, wherein the product ion having the second mass-to-charge ratio are product ions formed by charged loss of the fixed charge from the precursor ions.

4. The method of claim 1, further comprising the step of:

(6) determining the identity of the derivatized peptide or protein.

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5. The method of claim 4, wherein the step of determining the identity of the derivatized peptide or protein is performed by first repeating steps (1), (2), (3) and (4) and then subjecting the product ions having the second mass-to-charge ratio to dissociation to form a series of product ions having a range of mass-to-charge ratios, for the purpose of determining
30 the amino acid sequence of the peptide or protein.

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6. The method of claim 5, wherein the product ions having the second mass-to-charge ratio are formed by neutral loss from the precursor.
7. The method of claim 5, wherein the wherein the product ion having the second mass-
5 to-charge ratio is formed by charged loss from the precursor ion.
8. The method of claim 4, wherein the step of determining the identity of the derivatized peptide or protein is performed by use of high resolution mass analyzers.
- 10 9. The method of claim 8, wherein the use of high resolution mass analyzers provides mass accuracies of approximately 1-5 ppm on the product ion detected in step (5), or its complementary product ion.
10. The method of claim 4, wherein the step of determining the identity of the derivatized
15 peptide or protein comprises database searching to identify those peptides found to contain a fixed charge derivative.
11. The method of claim 1, wherein the step of dissociation comprises a method selected from the group consisting of (i) collisions with an inert gas (collision-induced dissociation
20 (CID) or collisionally-activated dissociation (CAD)); (ii) collisions with a surface (surface-induced dissociation (SID)); (iii) interaction with photons resulting in photodissociation, optionally using a laser; (iv) thermal/black body infrared radiative dissociation (BIRD); and (v) interaction with an electron beam, resulting in electron-induced dissociation for singly charged cations (EID), electron-capture dissociation (ECD) for multiply charged cations, or
25 combinations thereof.
12. The method of any of claims 1-11, wherein the method is used for identification of amino acids, peptides or proteins.
- 30 13. The method of any of claims 1-11, wherein the method is used for quantitation of amino acids, peptides or proteins.

14. The method of any of claims 1-11, wherein the method is used for amino acid, peptide or protein differential quantitation based on the incorporation of suitable isotopic or structural labels to the fixed charge.
- 5 15. The method of claim 14, wherein the isotopic labels are one or more of ^{13}C , ^{15}N , and ^2H .
16. The method of any of claims 1-11, wherein the method is used for analysis of post translational modification status of amino acids, peptides or proteins.
- 10 17. The method of claim 16, wherein the analysis of post translational modification status of amino acids, peptides or proteins comprises incorporation of the fixed-charge derivative via a β -elimination/Michael addition method for forming mass spectrometry stable derivatives of O-phosphorylated and O-glycosylated serine, or O-phosphorylated and O-
- 15 glycosylated threonine.
18. The method of any of claims 1-11, wherein the method is used for analysis of cross-linking status of amino acids, peptides or proteins.
- 20 19. The method of any of claims 1-11, wherein the method is used for analysis of interaction of proteins.
20. The method of any of claims 1-19, wherein the fixed-charge derivative is contained on the side-chain of a selected amino acid residue or a side-chain of a selected amino acid
- 25 residue contained within a protein or peptide.
21. The method of claim 20, wherein the selected amino acid residue is that of a rare amino acid.
- 30 22. The method of claim 20, wherein the selected amino acid residue contains a S atom in the side chain thereof.

23. The method of claim 22, wherein the amino acid residue is methionine, cysteine, homocysteine or selenocysteine.
24. The method of claim 23, wherein the amino acid residue is methionine and wherein
5 the method is performed according to any of Schemes 1, 3 and 4.
25. The method of claim 23, wherein the amino acid residue is cysteine and wherein the method is performed according to any of Schemes 2, 5, 6, 7 and 8.
- 10 26. The method of claim 20, wherein the selected amino acid residue is tryptophan or tyrosine.
27. The method of claim 26, wherein the amino acid residue is tyrosine or tryptophan and wherein the method is performed according to any of Schemes 5 and 6.
- 15 28. The method of claim 20, wherein the side chain contains an S-alkyl group.
29. The method of claim 28, wherein the amino acid residue is methionine, S-alkyl cysteine, S-alkyl homocysteine, S-alkyl tryptophan or S-alkyl tyrosine.
- 20 30. The method of any of claims 1-19, wherein the fixed-charge derivative is contained on a side-chain of a post-translationally modified amino acid residue.
31. The method of claim 30, wherein the fixed-charge derivative is contained on an O-
25 linked post-translationally modified amino acid residue.
32. The method of claim 30, wherein the O-linked post-translationally modified amino acid residue is a dehydroalanine residue formed by β -elimination from an O-linked post-translationally modified serine amino acid residue (Scheme 9).
- 30 33. The method of claim 30, wherein the O-linked post-translationally modified amino acid residue is a dehydroamino-2-butyric acid residue formed by β -elimination from an O-linked post-translationally modified threonine amino acid residue.

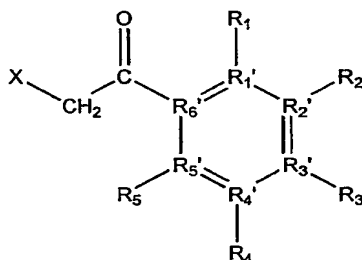
34. The method of any of claims 1-19, wherein the fixed-charge derivative is contained on a cross-link contained between two amino acids, peptides or proteins (Scheme 10).
- 5 35. The method of any of claims 1-19, wherein the fixed-charge derivative is contained within a cross-linking reagent.
36. The method of any of claims 1-35, wherein the fixed-charge derivative is selectively pre-enriched by solid phase capture methods using fixed charge reagents covalently coupled
10 to beads or insoluble polymers (Scheme 11).
37. The method of any of claims 1-36, wherein the fixed-charge ion is a sulfonium ion, a quaternary alkylammonium or a quaternary alkylphosphonium ion.
- 15 38. The method of any of claims 1-37, wherein the analysis of the amino acid, peptide or protein ion is performed by tandem mass spectrometry.
39. The method of claim 38, wherein the tandem mass spectrometer is equipped with electrospray ionization (ESI) or matrix assisted laser desorption ionization (MALDI)
20 interfaces to transfer the protein or peptide ion into the gas-phase.
40. The method of claim 38, wherein the tandem mass spectrometer is a tandem-in-space mass spectrometer, a tandem-in-time mass spectrometer, or a combination thereof.
- 25 41. The method of claim 40, wherein the tandem-in-space mass spectrometer is a sector mass spectrometer, a time of flight mass spectrometer, a triple quadrupole mass spectrometer, or a hybrid mass spectrometer combining time of flight and quadrupole instruments.
42. The method of claim 41, wherein the sector mass spectrometer is a double focusing
30 sector mass spectrometer or a hybrid mass spectrometer combining sector and quadrupole instruments.

43. The method of claim 38, wherein the tandem-in-time mass spectrometer is a two-dimensional quadrupole ion trap mass spectrometer, a three-dimensional quadrupole ion trap mass spectrometer or a Fourier-transform ion cyclotron resonance (FT-ICR) mass spectrometer.

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44. The method of any of the foregoing claims, wherein the method further includes one or more steps of protein extraction, protein separation, reduction and alkylation of cysteine disulfides and/or digestion.

10 45. The method of any of the foregoing claims, wherein the amino acids, peptides or proteins are derivatized using a substituted acetophenone, or a salt thereof, or a solvate thereof, having the following formula:



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46. The method of claim 45, wherein X is any halogen, sulfonic ester, perchlorate ester or chlorosulfonate.

47. The method of claim 45, wherein $\text{R}_1\text{-R}_5$ are H, and $\text{R}_1'\text{-R}_6'$ are ^{12}C .

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48. The method of claim 45, wherein the substituted acetophenone is an isotopically encoded substituted acetophenone, or a salt thereof, or a solvate thereof.

49. The method of claim 48, wherein X is any halogen, sulfonic ester, perchlorate ester or chlorosulfonate.

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50. The method of claim 49, wherein at least one of, and preferably at least three of $\text{R}_1\text{-R}_5$ are ^2H , and $\text{R}_1'\text{-R}_6'$ are ^{12}C .

51. The method of claim 49, wherein R_1 - R_5 are H, and at least one of, and preferably at least three of R_1' - R_6' are ^{13}C .

5 52. The method of any of claims 45-51, wherein at least one of R_1 - R_5 is a functional group containing an atom other than hydrogen or carbon.

53. The method of any of claims 45-52, wherein the substituted acetophenone is water soluble.

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54. The method of claim 53, wherein

R_1 is SO_2H , and R_2 - R_5 are H, and R_1' - R_6' are ^{12}C ;

R_1 is H, R_2 is SO_2H , and R_3 - R_5 are H, and R_1' - R_6' are ^{12}C ;

R_{1-2} are H, R_3 is SO_2H , and R_4 - R_5 are H, and R_1' - R_6' are ^{12}C ;

15 R_1 is SO_3H , and R_2 - R_5 are H, and R_1' - R_6' are ^{12}C ;

R_1 is H, R_2 is SO_3H , and R_3 - R_5 are H, and R_1' - R_6' are ^{12}C ; or

R_{1-2} are H, R_3 is SO_3H , and R_4 - R_5 are H, and R_1' - R_6' are ^{12}C .

55. The method of claim 53, wherein

20 R_1 is SO_2H , and at least one of, and preferably at least three of R_2 - R_5 are ^2H , and R_1' - R_6' are ^{12}C ;

R_1 is SO_2H , and R_2 - R_5 are H, and at least one of, and preferably at least three of R_1' - R_6' are ^{13}C ;

R_2 is SO_2H , and at least one of, and preferably at least three of R_1 and R_3 - R_5 are ^2H ,
25 and R_1' - R_6' are ^{12}C ;

R_2 is SO_2H , and R_1 and R_3 - R_5 are H, and at least one of, and preferably at least three of R_1' - R_6' are ^{13}C ;

R_3 is SO_2H , and at least one of, and preferably at least three of R_1 - R_2 and R_4 - R_5 are ^2H , and R_1' - R_6' are ^{12}C ;

30 R_3 is SO_2H , and R_1 - R_2 and R_4 - R_5 are H, and at least one of, and preferably at least three of R_1' - R_6' are ^{13}C ;

R_1 is SO_3H , and at least one of, and preferably at least three of R_2 - R_5 are ^2H , and R_1' - R_6' are ^{12}C ;

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R_1 is SO_3H , and R_2 - R_5 are H, and at least one of, and preferably at least three of R_1' - R_6' are ^{13}C ;

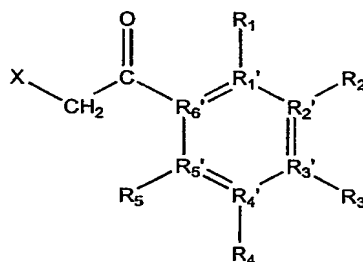
R_2 is SO_3H , and at least one of, and preferably at least three of R_1 and R_3 - R_5 are ^2H , and R_1' - R_6' are ^{12}C ;

5 R_2 is SO_3H , and R_1 and R_3 - R_5 are H, and at least one of, and preferably at least three of R_1' - R_6' are ^{13}C ;

R_3 is SO_3H , and at least one of, and preferably at least three of R_1 - R_2 and R_4 - R_5 are ^2H , and R_1' - R_6' are ^{12}C ; and

10 R_3 is SO_3H , and R_1 - R_2 and R_4 - R_5 are H, and at least one of, and preferably at least three of R_1' - R_6' are ^{13}C .

56. A substituted acetophenone, or a salt thereof, or a solvate thereof, having the following formula:



wherein X is a sulfonic ester, perchlorate ester or chlorosulfonate.

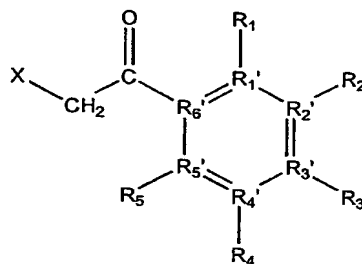
57. The substituted acetophenone of claim 56, wherein R_1 - R_5 are H, and R_1' - R_6' are ^{12}C .

58. The substituted acetophenone of claim 56, wherein the substituted acetophenone is an isotopically encoded substituted acetophenone, or a salt thereof, or a solvate thereof.

59. The substituted acetophenone of claim 58, wherein at least one of, and preferably at least three of R_1 - R_5 are ^2H , and R_1' - R_6' are ^{12}C .

60. The substituted acetophenone of claim 58, wherein R_1 - R_5 are H, and at least one of, and preferably at least three of R_1' - R_6' are ^{13}C .

61. A water soluble substituted acetophenone, or a salt thereof, or a solvate thereof, having the following formula:



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62. The water soluble substituted acetophenone of claim 61, wherein X is any halogen, sulfonic ester, perchlorate ester or chlorosulfonate.

10 63. The water soluble substituted acetophenone of claim 62, wherein X is Br or I.

64. The water soluble derivative of the substituted acetophenone of claim 63, wherein R₁ is SO₂H, R₂-R₅ are H, and R₁'-R₆' are ¹²C;

R₁ is H, R₂ is SO₂H, R₃-R₅ are H, and R₁'-R₆' are ¹²C;

15 R₁₋₂ are H, R₃ is SO₂H, R₄-R₅ are H, and R₁'-R₆' are ¹²C;

R₁ is SO₃H, R₂-R₅ are H, and R₁'-R₆' are ¹²C;

R₁ is H, R₂ is SO₃H, R₃-R₅ are H, and R₁'-R₆' are ¹²C; or

R₁₋₂ are H, R₃ is SO₃H, R₄-R₅ are H, and R₁'-R₆' are ¹²C.

20 65. An isotopically encoded form of the water soluble substituted acetophenone of claim 61.

66. The isotopically encoded water soluble substituted acetophenone of claim 64, wherein X is any halogen, sulfonic ester, perchlorate ester or chlorosulfonate.

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67. The isotopically encoded water soluble substituted acetophenone of claim 66, wherein R₁ is SO₂H, at least one of, and preferably at least three of R₂-R₅ are ²H, and R₁'-R₆' are ¹²C;

R_1 is SO_2H , R_2 - R_5 are H, and at least one of, and preferably at least three of R_1' - R_6' are ^{13}C ;

R_2 is SO_2H , at least one of, and preferably at least three of R_1 and R_3 - R_5 are ^2H , and R_1' - R_6' are ^{12}C ;

5 R_2 is SO_2H , R_1 and R_3 - R_5 are H, and at least one of, and preferably at least three of R_1' - R_6' are ^{13}C ;

R_3 is SO_2H , at least one of, and preferably at least three of R_1 - R_2 and R_4 - R_5 are ^2H , and R_1' - R_6' are ^{12}C ;

10 R_3 is SO_2H , R_1 - R_2 and R_4 - R_5 are H, and at least one of, and preferably at least three of R_1' - R_6' are ^{13}C ;

R_1 is SO_3H , at least one of, and preferably at least three of R_2 - R_5 are ^2H , and R_1' - R_6' are ^{12}C ;

R_1 is SO_3H , R_2 - R_5 are H, and at least one of, and preferably at least three of R_1' - R_6' are ^{13}C ;

15 R_2 is SO_3H , at least one of, and preferably at least three of R_1 and R_3 - R_5 are ^2H , and R_1' - R_6' are ^{12}C ;

R_2 is SO_3H , R_1 and R_3 - R_5 are H, and at least one of, and preferably at least three of R_1' - R_6' are ^{13}C ;

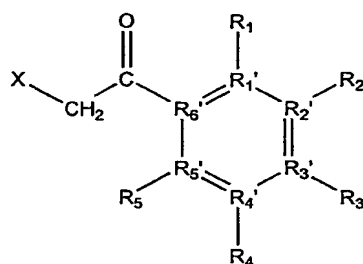
20 R_3 is SO_3H , at least one of, and preferably at least three of R_1 - R_2 and R_4 - R_5 are ^2H , and R_1' - R_6' are ^{12}C ;

R_3 is SO_3H , R_1 - R_2 and R_4 - R_5 are H, and at least one of, and preferably at least three of R_1' - R_6' are ^{13}C .

68. A reagent kit for analysis of amino acids, peptides or proteins by mass spectrometry
25 comprising a container containing the substituted acetophenone of any of claims 56-67.

69. A reagent kit for analysis of amino acids, peptides or proteins by mass spectrometry comprising a container containing a substituted acetophenone, or a salt thereof, or a solvate thereof, having the following formula:

- 73 -



wherein X is a halide.

- 5 70. The reagent kit of claim 69, wherein X is Br or I.
71. The reagent kit of claim 69 or 70, wherein R_1 - R_5 are H, and R_1' - R_6' are ^{12}C .
72. The reagent kit of claim 69, wherein the substituted acetophenone is an isotopically
10 encoded substituted acetophenone, or a salt thereof, or a solvate thereof.
73. The reagent kit of claim 72, wherein at least one of, and preferably at least three of R_1 - R_5 are ^2H , and R_1' - R_6' are ^{12}C .
- 15 74. The reagent kit of claim 72, wherein R_1 - R_5 are H, and at least one of, and preferably at least three of R_1' - R_6' are ^{13}C .
75. The reagent kit of claim 68-74, further comprising one or more containers containing:
cysteine disulfide reducing agents, cysteine alkylating reagents, proteases or chemical
20 cleavage agents, and solvents.
76. The reagent kit of claim 75, wherein the cysteine disulfide reducing agents are:
dithiothreitol (DTT), β -mercaptoethanol, tris-carboxyethyl phosphine (TCEP), and/or
tributylphosphine (TBP).
- 25 77. The reagent kit of claim 75, wherein the cysteine alkylating reagents are alkylhalides
(e.g. iodoacetic acid, iodoacetamide), vinylpyridine or acrylamide.

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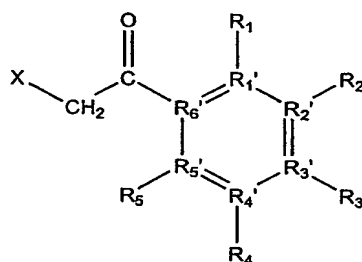
78. The reagent kit of claim 75, wherein the proteases or chemical cleavage agents are trypsin, Endoproteinase Lys-C, Endoproteinase Asp-N, Endoproteinase Glu-C, pepsin, papain, thermolysin, cyanogen bromide, hydroxylamine hydrochloride, 2-[2'-nitrophenylsulfenyl]-3-methyl-3'-bromoindole (BNPS-skatole), iodosobenzoic acid,
 5 pentafluoropropionic acid and/or dilute hydrochloric acid.

79. The reagent kit of claim 75, wherein the solvents are urea, guanidine hydrochloride, acetonitrile, methanol and/or water.

10 80. An amino acid or peptide comprising an amino acid derivatized to include a side chain fixed-charge sulfonium ion, quaternary alkylammonium ion or quaternary alkylphosphonium ion.

81. The amino acid or peptide of claim 80, wherein the amino acid is derivatized using
 15 the substituted acetophenone of any of claims 55-66.

82. The amino acid or peptide of claim 80, wherein the amino acid is derivatized using a substituted acetophenone, or a salt thereof, or a solvate thereof, having the following formula:



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wherein X is a halide.

83. The amino acid or peptide of claim 82, wherein X is Br or I.

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84. The amino acid or peptide of claim 82 or 83, wherein R_1 - R_5 are H, and R_1' - R_6' are ^{12}C .

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85. The amino acid or peptide of claim 82, wherein the substituted acetophenone is an isotopically encoded substituted acetophenone, or a salt thereof, or a solvate thereof.

86. The amino acid or peptide of claim 85, wherein at least one of, and preferably at least
5 three of R_1 - R_5 are ^2H , and R_1' - R_6' are ^{12}C .

87. The amino acid or peptide of claim 85, wherein R_1 - R_5 are H, and at least one of, and preferably at least three of R_1' - R_6' are ^{13}C .

10 88. The amino acid or peptide of any of claims 80-87, wherein the amino acid derivative is isotopically encoded.

89. A method for providing an internal standard in a mass spectrometer method comprising adding to a sample a predetermined quantity of the fixed charge derivatized
15 amino acid or peptide claimed in claim 80-88.